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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/687,558	10/15/2003	Glen S. Kwon	33-02	7151
23713 7590 08/23/2007 GREENLEE WINNER AND SULLIVAN P C 4875 PEARL EAST CIRCLE SUITE 200 BOULDER, CO 80301			EXAMINER KISHORE, GOLLAMUDI S	
			ART UNIT 1615	PAPER NUMBER
			MAIL DATE 08/23/2007	DELIVERY MODE PAPER

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

Office Action Summary	Application No. 10/687,558	Applicant(s) KWON, GLEN S.	
	Examiner Gollamudi S. Kishore, Ph.D	Art Unit 1615	

-- The MAILING DATE of this communication appears on the cover sheet with the correspondence address --

Period for Reply

A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 3 MONTH(S) OR THIRTY (30) DAYS, WHICHEVER IS LONGER, FROM THE MAILING DATE OF THIS COMMUNICATION.

- Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication.
- If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication.
- Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).

Status

- 1) ☒ Responsive to communication(s) filed on 25 July 2007.
- 2a) ☒ This action is **FINAL**. 2b) ☐ This action is non-final.
- 3) ☐ Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under *Ex parte Quayle*, 1935 C.D. 11, 453 O.G. 213.

Disposition of Claims

- 4) ☒ Claim(s) 1-15 is/are pending in the application.
- 4a) Of the above claim(s) _____ is/are withdrawn from consideration.
- 5) ☐ Claim(s) _____ is/are allowed.
- 6) ☒ Claim(s) 1-15 is/are rejected.
- 7) ☐ Claim(s) _____ is/are objected to.
- 8) ☐ Claim(s) _____ are subject to restriction and/or election requirement.

Application Papers

- 9) ☐ The specification is objected to by the Examiner.
- 10) ☐ The drawing(s) filed on _____ is/are: a) ☐ accepted or b) ☐ objected to by the Examiner.
Applicant may not request that any objection to the drawing(s) be held in abeyance. See 37 CFR 1.85(a).
Replacement drawing sheet(s) including the correction is required if the drawing(s) is objected to. See 37 CFR 1.121(d).
- 11) ☐ The oath or declaration is objected to by the Examiner. Note the attached Office Action or form PTO-152.

Priority under 35 U.S.C. § 119

- 12) ☐ Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).
- a) ☐ All b) ☐ Some * c) ☐ None of:
1. ☐ Certified copies of the priority documents have been received.
 2. ☐ Certified copies of the priority documents have been received in Application No. _____.
 3. ☐ Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)).

* See the attached detailed Office action for a list of the certified copies not received.

Attachment(s)

- | | |
|--|---|
| 1) <input type="checkbox"/> Notice of References Cited (PTO-892) | 4) <input type="checkbox"/> Interview Summary (PTO-413) |
| 2) <input type="checkbox"/> Notice of Draftsperson's Patent Drawing Review (PTO-948) | Paper No(s)/Mail Date. _____ |
| 3) <input type="checkbox"/> Information Disclosure Statement(s) (PTO/SB/08) | 5) <input type="checkbox"/> Notice of Informal Patent Application |
| Paper No(s)/Mail Date _____ | 6) <input type="checkbox"/> Other: _____ |

DETAILED ACTION

The amendment dated 7-25-07 is acknowledged.

Claims included in the prosecution are 1-15.

Upon consideration, the 112 rejection and the 103 rejection over Moribe et al are withdrawn.

Claim Rejections - 35 USC § 103

1. The following is a quotation of 35 U.S.C. 103(a) which forms the basis for all obviousness rejections set forth in this Office action:

(a) A patent may not be obtained though the invention is not identically disclosed or described as set forth in section 102 of this title, if the differences between the subject matter sought to be patented and the prior art are such that the subject matter as a whole would have been obvious at the time the invention was made to a person having ordinary skill in the art to which said subject matter pertains. Patentability shall not be negated by the manner in which the invention was made.

2. Claims 1-15 are rejected under 35 U.S.C. 103(a) as being unpatentable over Onyuksel et al (6,217,886) of record.

Onyuksel et al disclose a method of preparation of micelles containing polyene compounds, Amphotericin B and Nystatin. The method involves dissolving the water insoluble compound and the lipid conjugated to a Water-soluble polymer (PEG-DSPE) in an organic solvent, removal of the organic solvent and hydrating the lipid film to form micelles (col. 14, lines 15-47; claims 7-11 and 31). The composition further includes a cryopreservative (col. 14, line 67). What are lacking in Onyuksel et al are the pressure and temperature conditions under which the organic solvent is removed from the PEG-DSPE, active agent solution before hydrating it. However, in the absence of showing unexpected results, evaporation of a solvent is a manipulatable parameter in the highly

developed chemical sciences. The examiner also points out that Onyuksel et al in Example 1 teach the use of rotoevaporator to remove the solvent. Since rotoevaporation is done under vacuum conditions, the pressure is lower than the atmospheric pressure even possibly including the claimed pressures. Onyuksel et al also lacks the teachings of the ratios of PEG-DSPE to amphotericin B. In the examples, Onyuksel et al teach the amounts of the active agent in terms of weight and not moles. In the absence of showing the criticality, it is deemed obvious to manipulate the basic teachings of Onyuksel et al to obtain micelles with the desired amounts of the active agents. Finally it should be pointed out that Onyuksel et al specifically disclose dextrose as the cryopreservative. However, since dextrose is a sugar and sugars are known cryopreservatives, one of ordinary skill in the art at the time the invention was made would expect reasonable expectation of success using dextrose.

Applicant's arguments have been fully considered, but are not found to be persuasive. Applicant argues that Onyuksel does not disclose that a polyene antibiotic-containing antibiotic preparation (made by the method of Onyuksel) would contain deaggregated antibiotic (for example, Am B). According to applicant the present invention relies on deaggregated Am B as a means for minimizing hemolytic activity while providing a soluble formulation of this difficult-to-administer therapeutic agent and that the presently claimed preparation process allows for the association of the polyene antibiotic with PEG-DSPE in a way which prevents aggregation of antibiotic when hydrated, thus minimizing its toxicity (as measured by hemolysis). Further according to applicant, the cited Onyuksel actually teaches away from the present claimed method

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for providing a polyene antibiotic formulation which is soluble and has reduced toxicity (due to deaggregated state) as compared to known prior art compositions, either with respect to the polymer carrier or aggregated polyene antibiotic. These arguments are not persuasive. Claims 7-9 in Onyuksel teach the same method of preparation of the composition containing the biologically active agent and the compounds taught by Onyuksel are amphotericin B and Nystatin and therefore, one would expect similar product and applicant has not shown that amphotericin B in Onyuksel is not in a deaggregated form. Applicant argues that Onyuksel does not teach instant conditions of hydration. The examiner points out these are manipulatable parameters practiced by an artisan. Applicant argues that Onyuksel teaches a crystalline product or a sterically stabilized micelle preparation and instant invention does not relate to a crystalline product. This argument is not persuasive since as recognized by applicants themselves, Onyuksel teaches both crystalline and micellar products (hydrated). Claim 9 in Onyuksel clearly indicates micellar formation.

3. Claims 1-15 are rejected under 35 U.S.C. 103(a) as being unpatentable over Allen et al (US 2004/0013717) by itself or in view of Yu et al (Journal of Controlled Release, 1998) of record or vice versa.

Allen et al disclose micellar formulations containing PEG-DSPE to deliver any chemically or biologically active agent. The method of preparation involves dissolving the active agent and the phospholipid in an organic solvent, evaporation of the organic solvent using a rotary evaporator increasing the vacuum in increments of 25 mbar and hydrating the lipid film to form the micelles. The composition can be freeze-dried in the

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presence of a cryoprotectant such as a saccharide and rehydrated before use. The molecular weight of PEG is between 1000-10,000. (0016-0018, 0025, 0028, 0030, 0035, 0087, Examples 1 and 2). Although Allen et al do not specifically teach that the active agent to be amphotericin B, it would have been obvious to one of ordinary skill in the art to use any active agent including amphotericin B with a reasonable expectation of success since Allen et al teach its general applicability and provide guidance.

Yu et al teach polymeric micelles for the delivery of amphotericin. The polymer used for the formation of micelles is a PEG derivative of aspartic acid. According to Yu, the use of the polymeric micelles reduces the haemolytic activity of amphotericin B (abstract).

One of ordinary skill in the art would be motivated to use amphotericin B as an active agent in the micelles of Allen et al with a reasonable expectation of success since the reference of Yu et al shows the knowledge in the art of encapsulation of PEG containing polymeric micelles for the reduction of haemolytic activity of amphotericin B. Alternately, the use of PEG-DSPE instead of PEG-asp in the micelles of Yu et al would have been obvious to one of ordinary skill in the art since the reference of Allen et al shows that PEG-DSPE also forms micelles and such micelles could be used for the delivery of active agents. Although Allen et al do not specifically teach dextrose as the saccharide, the use of any saccharide would have been obvious to one of ordinary skill in the art with a reasonable expectation of success.

Applicant's arguments have been fully considered, but are not found to be persuasive. Applicant argues that Allen focuses on photosensitizers and suggests the

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use of any chemically or biologically active agent and that this represents a vastly broad class of molecules, of which the polyene antibiotics are a small class. This argument is not persuasive since as recognized by applicants themselves, Allen is suggestive of the use any active agent and therefore, it is within the skill of the art to use any active agent including polyene antibiotics with a reasonable expectation of success (see also Supreme court decision in *KSR International Co. V. Teleflex Inc.*, 550 U.S. -, 82 USPQ2d 1385 (2007)).

Applicant argues that Yu provides an alternative strategy to that claimed for formulating Am B. According to applicant, Yu teaches a different polymeric material and teaches a different method and that there is no indication that various components such as the polymer are interchangeable. This argument is not persuasive since Yu shows the knowledge in the art of encapsulating Am B in micellar formulations that too with a polymer containing micelles. Therefore, one of ordinary skill in the art would be motivated to use the micelles of Allen to encapsulate Am B. In response to applicant's argument that the examiner's conclusion of obviousness is based upon improper hindsight reasoning, it must be recognized that any judgment on obviousness is in a sense necessarily a reconstruction based upon hindsight reasoning. But so long as it takes into account only knowledge which was within the level of ordinary skill at the time the claimed invention was made, and does not include knowledge gleaned only from the applicant's disclosure, such a reconstruction is proper. See *In re McLaughlin*, 443 F.2d 1392, 170 USPQ 209 (CCPA 1971).

4. Claim 14 is rejected under 35 U.S.C. 103(a) as being unpatentable over Onyuksel et al (6,217,886). OR Allen et al (US 2004/0013717) by itself or in view of Yu et al (Journal of Controlled Release, 1998) of record or vice versa as set forth above, further in view of McShane (6,906,042).

The teachings of Onyuksel et al, Allen et al, Yu et al have been discussed above. What is lacking in these references is the teaching of the use of dextrose.

McShane while disclosing micellar formulations teaches that lyophilized micellar preparations can be rehydrated with dextrose solution, which is suitable for intravenous administrations (col. 12, lines 15-23). The use of dextrose in the micellar compositions of Onyuksel et al, Allen et al and Yu et al would have been obvious to one of ordinary skill in the art since such a micellar preparation is suitable for intravenous administration as taught by McShane.

Applicant's arguments have been fully considered, but are not persuasive. Applicant argues that McShane appears to relate to micelles having a very specific compound of the formula A and apart from the common use of the term, 'micelles' and the mention of dextrose, the disclosure of McShane is not relevant to the patentability of the presently claimed invention. This argument is not persuasive since McShane teaches the advantage of lyophilizing the micellar composition and the use of cryopreservatives; the advantage being to be able to dilute and prepare a solution suitable for intravenous administration. This advantage would be the same irrespective of what compound is used in the micelles and applicant has not shown any unexpected results by using dextrose in the compositions.

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5. Applicant's amendment necessitated the new ground(s) of rejection presented in this Office action. Accordingly, **THIS ACTION IS MADE FINAL**. See MPEP § 706.07(a). Applicant is reminded of the extension of time policy as set forth in 37 CFR 1.136(a).

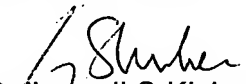
A shortened statutory period for reply to this final action is set to expire THREE MONTHS from the mailing date of this action. In the event a first reply is filed within TWO MONTHS of the mailing date of this final action and the advisory action is not mailed until after the end of the THREE-MONTH shortened statutory period, then the shortened statutory period will expire on the date the advisory action is mailed, and any extension fee pursuant to 37 CFR 1.136(a) will be calculated from the mailing date of the advisory action. In no event, however, will the statutory period for reply expire later than SIX MONTHS from the date of this final action.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Gollamudi S. Kishore, Ph.D whose telephone number is (571) 272-0598. The examiner can normally be reached on 6:30 AM- 4 PM, alternate Friday off.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Woodward Michael can be reached on (571) 272-8373. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

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Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see <http://pair-direct.uspto.gov>. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.


Gollamudi S Kishore, Ph.D
Primary Examiner
Art Unit 1615

GSK